

**REMARKS**

Claims 1 and 7 have been amended to specify that the inventive compositions are to ensure a retinoid half-life of at least about 20 days at 50 deg. C. Support for this amendment may be found in the Specification, particularly Example 6 on page 35. Additionally, Applicants are submitting together with this Amendment a Declaration from Dr. Iobst in support of the data and the claims.

New Claim 16 has been introduced to claim the booster alpha-ionone (B1) in combination with the booster(s) **AMEA or PAMEA** (fatty acid amide, B1 boosters), which in combination with the retinoid and the selected oils at POV of less than or equal to about 12. Support for this amendment is found throughout the Specification, and on pages 22 and 43 for example. Care has been taken not to introduce any new matter.

### The Present Invention

The present invention is directed to a combination of specified retinoids and specified retinoid boosters that are stabilized in a composition, wherein each constituent of the oil phase of the oil-in water emulsion has a peroxide value (POV) of less than or equal to about 12, preferably less than or equal to about 6. Controlling the POV is particularly important with combination of retinoids and boosters, to ensure a retinoid half-life of at least 20 days at 50 deg. C.

The present invention provides the dual benefit of enhancing retinoid efficacy within the skin while increasing the *stability of the retinoids in the composition* by the removal of any starting materials having a POV of greater than 12, and preferably greater than 6. See Specification at page 32.

The POV specification is critical for removing oil impurities that would have POV of greater than 12 and would thereby contribute to retinoid instability. For example, while the general category of mineral oil is known, depending on the commercial source of the oil, it may have undesirable impurities with POV greater than 12. Specifying the POV in the present claims excludes these impurities, or removes oils with POV greater than 12, which would promote instability. The resulting inventive compositions extend retinoid stability. Support for "removal of any starting materials having a peroxide value of greater than 12 and preferably greater than 6 may be found on page 32 and elsewhere in the Specification.

***Claims 1-2, 7-8, 13 and 15 Are Not Obvious Under 35 USC § 103***

Claims 1-2, 7-8, 13 and 15 were rejected under 35 U.S.C. 103(a) as being unpatentable over Granger et al. (USPN 5,716,627) or, in the alternative, in view of Potter et al. (US 5,620,692).

According to the Office Action, Granger et al. teaches a skin-conditioning composition comprising (a) retinol or retinyl ester, (b) azole (e.g. climbazole), (c) LMEA, and (d) cosmetically acceptable vehicle. The Office Action admits that *Granger '627 does not teach that each ingredient in the oil phase should have POV as recited in the claims*, i.e. that each constituent of the oil phase of the oil-in water emulsion has a peroxide value of less than or equal to about 12, preferably less than or equal to about 6. As Granger et al fail to disclose or suggest a POV, Potter et al. are cited. However, Applicants respectfully submit that Potter et al. fail to remedy the deficiencies of Granger '627. As such, a *prima facie* case of obviousness has not been made out.

Potter et al. fail to remedy the deficiencies of Granger '627 for the following reasons. While Potter et al. may define the POV parameter in general, it fails to teach or suggest specific materials and/or specific POV values as presently claimed. Potter et al. relate to a process to prepare oat oil compositions. See Abstract. The antioxidants in oat oil are esters of caffeic and ferulic acid, see Col. 1, lines 45-51, which have nothing to do with the oils claimed in the present invention and are antioxidants for skin lipids, while the low POV oils claimed herein have a stabilizing effect on other components of the cosmetic composition.

A composition with the same ingredients is not disclosed in Granger '627. The perfume in the Examples is not specified and therefore it does not specify all oil

components with a POV of less than or equal to 12. Applicants respectfully submit that, not all the named ingredients in the cited are the same, and they will not "invariably" possess the same characteristics. For example, while the general category of mineral oil is known, depending on the commercial source of the oil, it may have undesirable impurities with POV greater than 12. Specifying the POV in the present claims excludes these impurities, or removes oils with POV greater than 12, which would promote instability. The resulting inventive compositions extend *retinoid stability, defined by the concentration of retinoids in their original chemical form after a defined storage duration and temperature*. See present Specification at page 32.

Applicants are submitting together with this Amendment a Declaration by Dr. Lobst in support of objective evidence of stability of the inventive compositions as claimed. The data show that controlling the POV of the oil phase at below about 12 achieves a half-life of at least about 20 days at 50 deg. C.

**Claim 14, Requiring a Combination of LAMEA and alpha-Ionone, Is Free of the Art**

Claim 14 was rejected under 35 U.S.C. 103(a) as being unpatentable over Granger et al. (USPN 5,716,627) and Potter et al. as applied to 1-2, 7-8, 13 and 15, and further in view of Granger et al. (WO 98/13020) (to remedy lack of disclosure of alpha-ionone in the primary references).

Claim 14 requires the presence of the booster alpha-ionone (B1 booster) in combination with the booster LAMEA, which in combination with the retinoid and the selected oils at POV of less than or equal to about 12, are free of art.

Granger et al. (WO 98/13020) teaches away from using a fatty acid amide. *Evidence of teaching away is in the reference itself. Granger '020 in the Summary, on page 3 and in Claim 1 on pp. 52 and 54, state: "compound is not a fatty acid amide ...".* Therefore, a retinoid composition comprising LAMEA (B1 booster, fatty acid amide) would not be within the scope of Granger WO'020, and it would not be proper to combine it with Granger '627 and Potter. Accordingly, Claim 14, as amended, is in condition for allowance.

An obviousness rejection is proper only when "the subject matter as a whole would have been obvious at the time the invention was made ..." (emphasis added). 35 U.S.C. 103. Applicants respectfully submit that the Office Action has improperly chosen certain aspects of one reference and combined them with aspects of other references, without showing where the motivation is to combine them to come up with the subject matter of the present invention as a whole, within the meaning of 35 U.S.C. 103. Applicants submit that the pending claims are not obvious over the cited references, under 35 U.S.C. 103, especially in view of the present Amendment. Reconsideration and withdrawal of the rejection is respectfully requested.

Similarly, new Claim 16 is believed to be in condition for allowance.

**Conclusion**

The Granger et al. references and Potter et al., either alone or in combination, do not address the problem to which the present invention is addressed, i.e., improvements in stability of retinoids achieved by controlling POV of each constituent of the oil phase of the oil-in-water emulsion, while increasing the effectiveness of the retinoids.

Those skilled in the art would not have identified among the various materials a special stabilizing effect of oils with POV of less than about 12 when used in a formulation with retinoids and retinoid boosters from a mere reading of the cited art.

In view of the foregoing amendments and comments, Applicants request the Examiner to reconsider the rejections and now allow the claims.

If a telephone conversation would be of assistance, Applicant's undersigned attorney invites the Examiner to telephone at the number provided.

Respectfully submitted,

  
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